

K993092 SONADA ULTRASOUND SYSTEMApr 14, 2000
211 days to decisionK993092 · Product code: **OXO** · Radiology
Source: <https://www.510kdatabase.net/k993092/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Image-intensified Fluoroscopic X-ray System, Mobile (OXO)
Date received	Sep 16, 1999
Decision date	Apr 14, 2000
Days to decision	211 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	Sonovision Corp.
Location	Hasbrouck Heights, NJ, US
Contact	PAUL SCHNEIDER
510(k) history	1 submissions · 1 cleared · 2000-2000

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k993092/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated May 19, 2026